

L 8593-66

ACCESSION NR: AP5019896

ASSOCIATION: None

SUBMITTED: 08Apr65

ENCL: 01

SUB CODE: SS, EM

NR REF Sov: 004

OTHER: 004

Card 2/3

L 8593-66  
ACCESSION NR: AP5019896

ENCLOSURE: 01

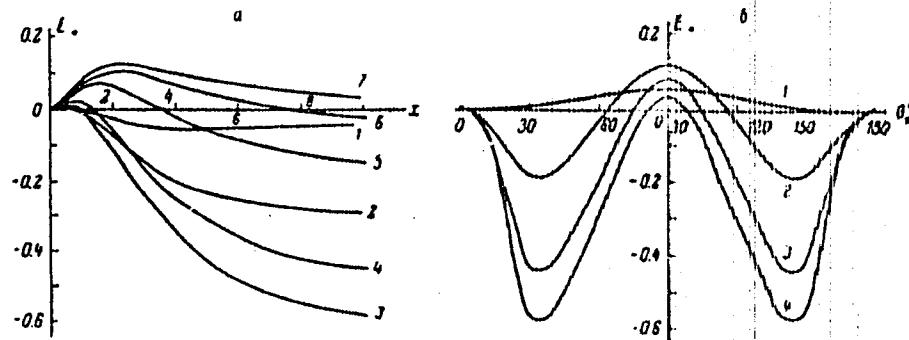


Fig. 1. Dependence of photomagnetic effect (arbitrary units) on  $x$  (left) and on the angle  $\theta$  (right) for different values of the angle and of  $x$ .

jw  
Card 3/3

L 36385-66 SMT(1) IJF(c)  
ACC NR: AP6014052

SOURCE CODE: UR/0056/66/050/004/1107/1123

53

AUTHOR: Kagan, Yu.; Zhernov, A. P.

ORG: none

TITLE: Theory of the electric conductivity of metals with nonmagnetic impurities

SOURCE: Zhurnal eksperimental'noy i teoreticheskoy fiziki, v. 50, no. 4, 1966,  
1107-1123

TOPIC TAGS: electric conductivity, atom, electron scattering, phonon spectrum,  
temperature dependence, metal impurity, impurity scattering

ABSTRACT: A theory is developed for the electric conductivity of metals with impurities. It consistently takes into account the deformation of the phonon spectrum occurring with the introduction of impurity atoms and also the arbitrary variations in the electron scattering amplitude on a separate ion. The impurity part of the resistance for the complete temperature range was found. It was shown that in the low temperature range, the electron scattering on the oscillating impurity ion leads to the appearance of the term  $\nu T^2$ . Interference between scattering on an impurity ion and on a perturbed phonon spectrum leads to the term  $\nu T^4$  and scattering on a deformed phonon spectrum leads to the term  $\nu T^5$ . At low impurity concentrations all these terms are proportional to the concentration. Greater temperatures of the impurity resistance produce a number of anomalies, especially in the case of heavy impurity atoms when a quasi-local level appears. At high temperatures, the impurity

Cord 1/2

L 22130-66 EWT(1) IJP(c) WW/QJ  
ACC NR: AF6004945

SOURCE CODE: UN/0056/65/050/001/0271/0280

AUTHOR: Kagan, Yu.

ORG: none

TITLE: Change in the resonance nuclear parameters during scattering by regular systems

SOURCE: Zhurnal eksperimental'noy i teoreticheskoy fiziki, v. 50, no. 1, 1966,  
271-280

TOPIC TAGS: nuclear resonance, nuclear scattering, compound nucleus, nuclear energy level, resonance line, line width, crystal lattice, elastic scattering, inelastic scattering

ABSTRACT: This is a sequel to an earlier investigation (ZhETF, Pis'ma v. 2, 130, 1965), dealing with the case of scattering by an isolated nucleus. Since the resonance parameters can be significantly altered in scattering from regular systems of identical nuclei, especially in the case when it is impossible to point to the individual nucleus responsible for the scattering, and the excitation of the resonance level or the formation of the compound nucleus have a collective character, the article is restricted to the case when the nuclei in the system are assumed

Card 1/2

L 22130-66

ACC NR: AP6004945

to have low lying resonance levels and to comprise a one-dimensional chain of identical nuclei. It is shown that if the nuclei form a regular one-dimensional chain or a two-dimensional array, then the elastic width and the position of the resonance level are noticeably changed. The change in the width is connected with the lifetime of the collective excitation of the state of the system of nuclei. The lifetime of such a state, and accordingly the width, can be either larger or smaller than the lifetime (width) of the isolated excited state. In the case of a three-dimensional crystal, the elastic width disappears completely and the energy dependence of the resonance interaction is determined only by the inelastic width. In the presence of spin in the ground state of the nucleus, and when an appreciable role is played by inelastic phonon scattering, the width takes an intermediate value between the inelastic width and the total width. The authors are grateful to A. I. Baz' for a valuable discussion. Orig. art. has: 2 figures and 36 formulas.

SUB CODE: 20/ SUBM DATE: 13Aug65/ ORIG REF: 004/ CTR REF: 002

Card 2/2 BK

L 116925-66 EWT(1)/EWT(m)/T/EWP(t)/ETI IJP(c) JD  
ACC NR: AP6015457 SOURCE CODE: UR/0101/66/008/005/1402/1416

AUTHOR: Brovman, Ye. G.; Kagan, Yu.

39  
B

ORG: none

TITLE: Phonon spectrum of the white tin lattice

SOURCE: Fizika tverdogo tela, v. 8, no. 5, 1966, 1402-1416

TOPIC TAGS: tin, phonon spectrum, Mossbauer effect

ABSTRACT: A systematic study is made of the dynamic problem of lattice oscillations within the framework of the Born-Karman model. Use is made of all force constants in the complete dynamic oscillation matrix for the first three coordination spheres; it is assumed that the interaction with the 4th, 5th, and 6th coordination spheres can be approximated for each sphere by one central and one noncentral force constant. Dispersion curves along three symmetric directions, the function phonon spectrum frequency distribution, heat capacity and Debye temperature, and probability of the Mossbauer effect are in good agreement with experimental data. Values of heat capacity at  $T > 100^{\circ}\text{K}$  and of probability of Mossbauer effect at  $T > 200^{\circ}\text{K}$  higher than those calculated are explained by anharmonism. Theoretical curves of anisotropy of Mossbauer effect compares favorably with experimental data (errors are indicated); value of anisotropy

Card 1/2

KAGAN, J.

PA

Synthesis of olefins and paraffinic hydrocarbons of low  
structure, containing a quaternary carbon atom. II  
Reaction between the hydrobromide of 1,1,3-trimethyl-  
butadiene and alkylmagnesium halides. R. Ya. Levin  
and Yu. B. Kagan. *J. Gen. Chem. (U. S. S. R.)* 11,  
523-6 (1941).—On the basis of previous work (C. A. 35,  
5002\*) the authors conclude that the hydrobromide of  
1,1,3-trimethylbutadiene (I) is 1,1,3-tetramethylallyl  
bromide (II), i.e., a tertiary bromide readily reactive with  
RMgX. In order to confirm this structure of I, its re-  
action with MeMgI was investigated. The reaction was  
carried out analogously to the previously reported reac-  
tion with cyclohexylmagnesium chloride. A 30% yield of a  
hydrocarbon (III) was isolated, b.p. 102-4°, nD<sup>20</sup> 1.4130,  
d<sup>20</sup> 0.7191; these const. are close to the known values  
for 2,4,4-trimethyl-2-pentene (IV). Ozonization of III in  
CHCl<sub>3</sub> yielded acetone (isolated as the peroxide) and  
Me<sub>2</sub>CO<sub>2</sub>H, thus further confirming the identity of III  
and IV. Catalytic hydrogenation of III yielded 2,4,4-  
trimethylpentane, which was identified by the phys.  
const. and the Raman spectrum. The information thus  
collected proves the identity of I and II. The reaction  
between I and RMgX should be a convenient prepn.  
method for saturated unsatd. hydrocarbons of low structure  
and const., a quaternary carbon atom; this is especially true  
for relatively little studied isoparaffins, Me<sub>2</sub>C(CH<sub>2</sub>)<sub>2</sub>Me<sub>2</sub>.  
By this reaction between I and EtMgBr there was prepd.  
2,4,4-trimethyl-2-hexene, b.p. 132°, nD<sup>20</sup> 1.4284, d<sup>20</sup> 0.7437;  
its hydrogenation (platinized charcoal) gave 2,4,4-tri-  
methylhexane, b.p. 129-30°, nD<sup>20</sup> 1.4081, d<sup>20</sup> 0.7306. Re-  
action of I with PrMgBr gave 2,4,4-trimethyl-2-heptene,  
b.p. 152.5-153.5°, nD<sup>20</sup> 1.4307, d<sup>20</sup> 0.7500. Hydrogenation  
of the latter yielded 2,4,4-trimethylheptane, b.p. 151-2°,  
nD<sup>20</sup> 1.4143, d<sup>20</sup> 0.7340. G. M. Knudsen

10

COMMON ELEMENTS

MATTEALS

ASM-SLA METALLURGICAL LITERATURE CLASSIFICATION

10000 10000 10000 10000

KASHAN, YIL  
B

27

\*812. Synthesis of  $\alpha,\beta$ -Dichlorethyl-Alkyl Ethers and Their Transformations. (In Russian.) M. F. Shogakovskii, Yu. B. Kugan, and F. P. Sidel'kovskaya, Journal of General Chemistry (U.S.S.R.), v. 17(10), May 1947, p. 967-968.

Conditions are described for the chlorination of vinyl ether in order to obtain dichlorethyl alkyl ethers. Methods are worked out for the preparation of chloroacetals by chlorination of vinyl ether in the presence of water or in a mixture with butyl alcohol. 16 ref.

AMSLA METALLURGICAL LITERATURE CLASSIFICATION

*Off Chalkley*

Use of the differential thermocouple for kinetic measurements. Yu. B. Kagan and A. N. Bashkirov (Inst. Combustible Minerals Acad. Sci. U.S.S.R., Moscow). *Izvest. Akad. Nauk S.S.R., Otdel. Tekh. Nauk* 1948, 349-48.

(1) On the assumption that the temp. difference between a catalyst on which a chem. reaction is taking place, and an inert substance placed in the same tube, is proportional to the rate of evolution (or absorption) of heat of reaction, hence proportional to the rate of the reaction, recording of the deflections  $\varphi$  of the galvanometer connected with the differential thermocouple (Ialandin and Patrikeev, C.A. 39, 8849) supplies a method of detn. of the fraction velocity  $dx/dt$ , provided the proportionality factor  $\varphi$  in  $\varphi = \varphi (dx/dt)$  is sufficiently independent of the temp. Fulfilment of the latter condition, involving practical temp. independence of the heat of reaction, of the sum of heat capacities, and of the heat conductance, was tested experimentally on the oxidation of Cu on a Kieselguhr carrier at 245° and at 80°; whereas at 245° oxidation is complete, its degree at 80°, calcd. on the assumption  $\varphi = \text{const.}$  (between 80 and 245°), was 27.37% as against 26.79% by analytical data, consequently, the assumption  $\varphi = \text{const.}$  is permissible. The general procedure for kinetic measurements involving a solid consists in plotting  $\varphi$  against the time  $t$ ; the area limited by the axis of abscissas, the curve and the ordinates  $\varphi_1$  and  $\varphi_2$  (at times  $t_1$  and  $t_2$ ) =  $\varphi (x_1 - x_1)$  where  $x$  = amt. of solid reacted. Integration from  $t = 0$  to  $t$  gives the area  $\varphi t$ . The concn. of the gas in the reaction zone being const., the rate of reaction (unless limited by the penetration of O<sub>2</sub> through a surface oxide film)  $dx/dt = k (a - x)$ , or  $\varphi = k \varphi (a - x)$ . In the case of a first-order reaction, the plot of  $\varphi$  against  $\varphi t$  is a straight line; its slope gives the rate const.  $k$ , the base of the triangle is =  $\varphi a$  ( $a$  = initial amt.). The error

due to loss of heat is immaterial provided the temp. difference does not exceed that for which Newton's law of cooling is valid, and the total heat capacity, heat cond., and the heat of reaction are practically const. in the given temp. range. (2) Expts. were made on Cu prep'd. by mixing 100 ml. kieselguhr with 2.5 g. Cu (as salt) in 75 ml. H<sub>2</sub>O, ppts. with KOH at 100°, washing with hot H<sub>2</sub>O, drying and granulating to cylinders 2-3 mm. long, and heating 1.5 hr. at 700°; the thermocouple was disposed between 5 ml. of the catalyst and 6 ml. of glass cylinders, sep'd. by mica, placed in the same tube of 2 cm. diam. The catalyst was reduced to Cu by H<sub>2</sub> at 6.5 l./hr. at 245°. Oxidation at 245° with a dry gas mixt. of 2.5% O<sub>2</sub> + 97.5% N<sub>2</sub> at 0.1 l./hr., was complete in 46 min.; further passage of air caused no deflection of the galvanometer. On successive oxidations of the same catalyst, following repeated reductions, the integrals corresponding to  $\varphi t$  increased regularly (e.g. 197, 210, 234 sq. cm.), indicating increasing activity of the Cu catalyst. At 80°, in an air stream at 0.1 l./hr., the reaction came to a halt after 11 min., further passage of air causing no change of  $\varphi$ . From the total area  $\varphi t$ , read on the  $\varphi (t)$  plot, as compared with the area corresponding to 100% oxidation (at 245°), the degree of oxidation at 80° was 27.37% of the Cu present (that oxidized at 245°). (3) By the linearity of the plot of  $\varphi$  against the  $\varphi t$  detd. by readings at various stages, the reaction at 80° is of the first order, with  $k = 0.08$  (time in min.). N. Then

KAGAN, YU. B.

USSR/Chemistry - Esters, Polymerization of  
Chemistry - Polymerization

Aug 48

"Significance of the Purity of Vinylalkyl Ethers on the Polymerisation Process:  
Polymerization of Vinylbutyl Ether in the Presence of Oxygen-Containing Organic  
Compounds," M. F. Shostakovskiy, F. P. Sidel'kovskaya, Yu. B. Kagan, Inst Org  
Chem, Acad Sci USSR, Lab of Vinyl Compounds, 6 $\frac{1}{2}$  pp

"Zhur Obshch Khimii" Vol XVIII (LXXX), No 8

The degree of polymerization of vinylalkyl esters, and, other things equal, the process of polymerization, depends mainly on purity of original ether. Oxygen-containing compounds -- alcohols, aldehydes, ketones and acetals -- found in commercial vinylalkyl ethers give rise to peroxide effect and reduce degree of polymerization of polyvinylalkyl ether formed. Mechanism of breaking of polymerization chains by alcohols is combination with initial vinylalkyl ethers or their polymers and formation of corresponding acetals. Submitted 20 Dec 45.

PA 19/49T27

KAGAN, Yu. B.

USSR/Chemistry - Synthesis  
Chemistry - Bonds

Jnn 49

"Synthesis of Aliphatic and Paraffin Hydrocarbons: VI, Diene Hydrocarbons in the  
Synthesis of Olefin Hydrocarbons With the Double Bond in the Center," R. Ya. Levina,  
V. A. Skvarchenko, Yu. B. Kagan, Ye. G. Treshchova, Lab of Org Chem imeni N. D. Zelinskiy,  
Moscow State Ord of Lenin U, 6 1/2 pp

"Zhur Obshch Khim" Vol XIX, No 1

In the reaction between allyl halides and organomagnesium compounds used for preparing 1-alkenes, the double bond causes an exchange of the halogen for a radical. In present study, 1,4-dichloro-2-butene and its bromine and analogue, prepared by halogenation of 1,3-butadiene, reacted with methylmagnesium bromide or iodide. Both halogens of these compounds were substituted by methyls theoretically forming 10% 3-hexane in the case of dibromide and up to 40% in the case of dichloride; 1,3-butadiene was formed as a by-product. The 1,2-dichloride of butadiene undergoes an allyl rearrangement to a 1,4-compound, and subsequently forms an alkene with the double bond in central position.

Submitted 29 Oct 47

58/49T42

KADAM, Yu. N.

26933. BASHKIROV, A. N., KOTLUK V. M. P., KADAM, Yu. N. - Kyopress i mekhanizmy sinteza  
li-tlevodoro dov iz okiri uleroda i vodoroda. Doklady akad nauk SSSR, 1949  
seriya, t. LVI, No 6 1949, S. 1029-31.

SO: "etopis' Zhurnal'nykh Statej", Vol. 36, 1949

*KAGAN, Yu. B.*

the mechanism of the synthesis of hydroquinone from carbon monoxide and hydrogen. *Zhurnal fizicheskoy kemi*, V. 48, No. 10, p. 2147-2151, 1973.

B. Kryukov, Yu. B. Kagan, and I. V. Khodatkin. *Zhurnal fizicheskoy kemi*, V. 48, No. 10, p. 2152-2156, 1973.

were 27.6, 27.7, 27.8, 28.0, 28.2, 28.4, 28.6, 28.8, 29.0, 29.2, 29.4, 29.6, 29.8, 30.0, 30.2, 30.4, 30.6, 30.8, 31.0, 31.2, 31.4, 31.6, 31.8, 32.0. Since  $\text{H}_2\text{O}$  is not observed as a product on the catalysts that catalyze (VIII) or reaction (III), a greater rate or equal to the velocity  $V_{\text{cat}}$  of reaction (III). The velocities of these reactions were studied directly on the catalysts selected. It appears that the reaction between the benzene and carbon monoxide could be interpreted as follows:

Further evidence is presented that the reaction of benzene with carbon monoxide over the catalysts studied is an addition reaction.

(100:10:5 245.6:100) - Contraction depressions (% found)

C19

Mechanism of the synthesis of hydrocarbons from carbon monoxide and hydrogen. A. N. Bashkinov, V. I. Kryukov, and Yu. B. Kryukov. Doklady Akad. Nauk SSSR, 178, 275-6 (1961).—The assumption (cf. C.A. 43, 44164) that the primary act in the Fischer-Tropsch synthesis is  $\text{CO} + 2\text{H}_2 \rightarrow \text{CH}_3 + \text{H}_2\text{O}$ , and not  $2\text{CO} + \text{H}_2 \rightarrow \text{CH}_4 + \text{CO}_2$ , and that  $\text{CO}_2$  is formed only as a result of the secondary reaction  $\text{CO} + \text{H}_2\text{O} \rightarrow \text{CO}_2 + \text{H}_2$ , was confirmed directly by flow expts. in which a very short contact time prevented the secondary reaction from taking place to any appreciable extent in one single pass, and the  $\text{H}_2\text{O}$  formed in each pass was removed by condensation between passes; in repeated recirculation. The expts. were conducted with a  $\text{CO:H}_2 = 1:2$  mixt. at  $300^\circ$  under 20 atm., at a gassing space velocity of 100,000 l./l. catalyst/hr.; practically complete conversion was ensured by repeated recycling, with the fresh gas admitted at the rate of 44 l./hr. Under these conditions, 200 l. gas gave, after several hrs., liquid hydrocarbons 33.6 g./cu. m., gaseous hydrocarbons ( $\text{C}_1 + \text{C}_2$ ) 41.2,  $\text{CH}_4$  100.0,  $\text{H}_2\text{O}$  249.3 g./cu. m., and no  $\text{CO}_2$ . This result excludes the direct reaction  $2\text{CO} + \text{H}_2 \rightarrow \text{CH}_4 + \text{CO}_2$ , and demonstrates that on Co, Ni, and Fe catalysts alike,  $\text{CO}_2$  is formed only through a secondary reaction between  $\text{CO}$  and  $\text{H}_2$ .

N. Tion

KAGAN, E. S.

Chemical Abst.  
Vol. 48 No. 9  
May 10, 1954  
Organic Chemistry

Role of oxygen-containing compounds in the synthesis of hydrocarbons from carbon monoxide and hydrogen. Yu. B. Kagan, Yu. B. Kravkov, M. V. Kaputkins, and A. N. Bushkinov. Bull. Acad. Sci. U.S.S.R., Div. Chem. Sci. 1952, 601-8 (Engl. translation). See C.A. 47, 3216. H. L. H.

KAGAN, Yu. B.

USSR/Chemistry - Synthetic Liquid Fuels Jul/Aug 52

"The Role of Oxygen-Containing Compounds in the Synthesis of Hydrocarbons From Carbon Monoxide and Hydrogen," Yu.-B. Kagan, Yu. B. Kryukov, Ye. V. Kamzolkina, A. N. Bashkirov, Petroleum Inst., Acad. Sci. USSR

"IZ Ak Nauk SSSR, Otdel Khim Nauk" No 4, pp 649-657

Article states that results of the expts described show that alcs cannot be regarded as intermediate products in the synthesis of hydrocarbons, and that iron catalysts, under the conditions of hydrocarbon synthesis, accelerate the oxidation of alcs and aldehydes. Advances hypotheses explaining the formation of oxygen-contg compds (by-products of hydrocarbons).

22916

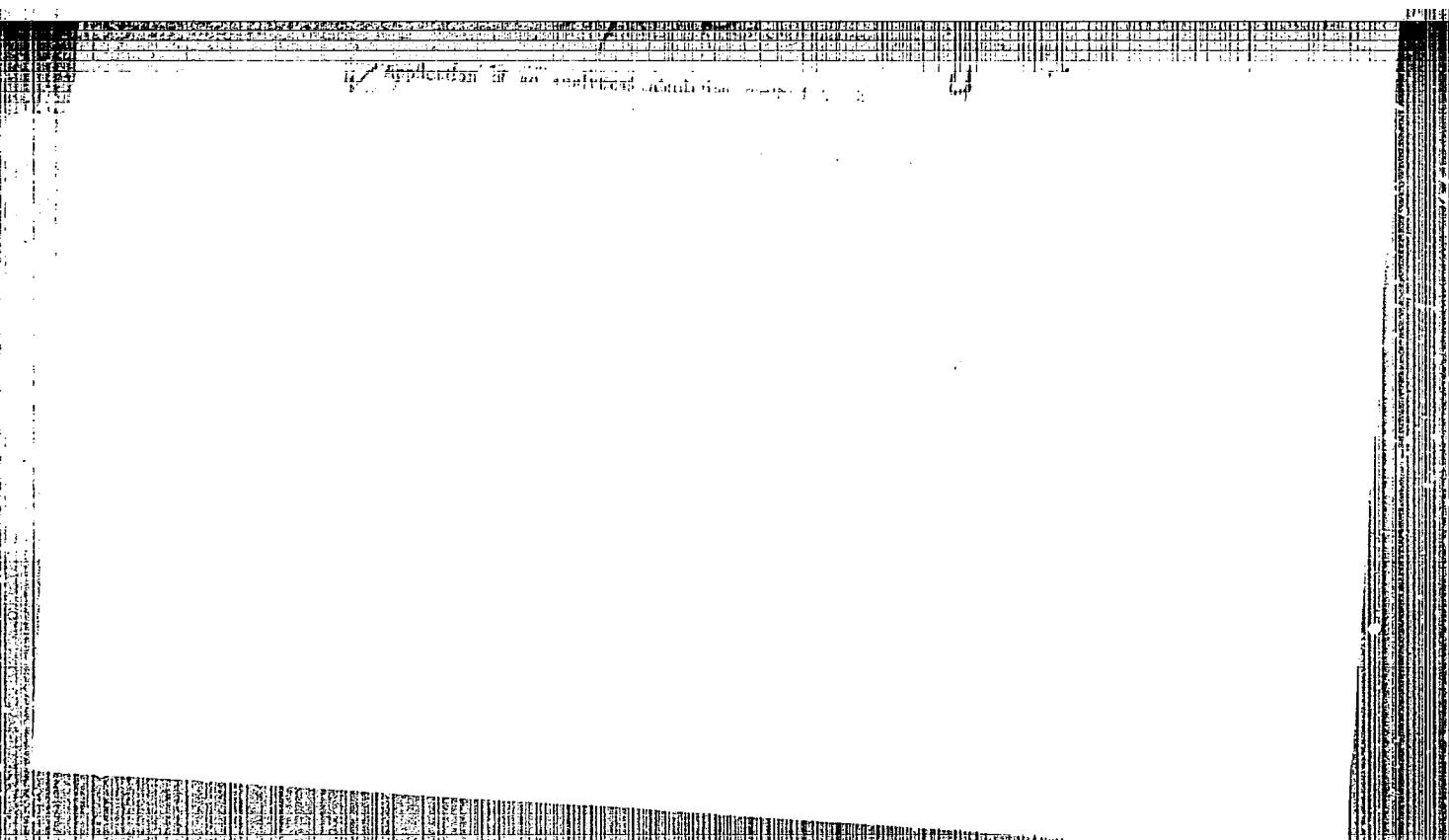
22916

KAGAN, Yu. R.

Oxidation of iron catalysts by water vapor in hydrourethane synthesis. A. N. Belyakov, Yu. B. Kagan, V. I. Kirillov, B. M. Fedorovich, and M. T. Dzhingarhan. *Zhur. Inst. Nefti Akad. Nauk S.S.R.*, 4, 151-8 (1957). Fe catalysts gradually lose their activity by the action of water vapor formed during the synthesis of hydrocarbons from CO and H (ratio 1:1) at 290° and av. pressure. The rate of Fe oxidation depends on the oxidation and reduction reactions which take place during the synthesis. Inactivation of Fe catalysts is due to the formation of FeO.  
M. Chaghmangarhan.

"APPROVED FOR RELEASE: 08/10/2001

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Knowing full well,

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APPROVED FOR RELEASE: 08/10/2001

CIA-RDP86-00513R000619910019-5"

BASHKIROV, A.N.; KAGAN, Yu.B.; LOKTEV, S.M.; MOROZOV, N.G.

Use of iron ore catalysts in the synthesis based on carbon monoxide and hydrogen. Trudy inst. nefti. 10:234-246 '57.  
(MIRA 11:4)

(Catalysts) (Hydrocarbons)

BASHKIROV, A.N.; KAGAN, Yu.B.; KOKTEV, S.M.; SHCHEKIN, V.V.; GOL'DIN, S.A.;  
MOROZOV, N.G.

Activating characteristics of molten iron catalysts used in the  
synthesis based on carbon monoxide and hydrogen, and reduced at  
high temperatures. Trudy inst. nefti. 10:247-261 '57.

(MIRA 11:4)

(Catalysts) (Hydrocarbons)

KAGAN, Yu.B.; BASHKIROV, A.N.; ZVEZDKINA, L.I.; ORLOVA, N.A.; KLIGER, G.A.

Influence of reduction conditions on the properties of molten  
iron catalysts used in alcohol synthesis from carbon monoxide  
and hydrogen. Trudy inst. nefti. 10:262-268 '57. (MIRA 11:4)  
(Alcohols) (Carbon monoxide) (Hydrogen)

KAGAN, Yu.B.; BASHKIROV, A.N.; KRYUKOV, Yu.B.; LOKTEV, S.M.

Formation of the active surface of fused iron catalysts for  
synthesis from CO and H<sub>2</sub>. Khim i tekhn. topl. i nasej } no.3:  
14-22 Mr '58. (MIRA 11:3)

1. Institut nefti AN SSSR.  
(Catalysts) (Iron oxides) (Hydrocarbons)

KHL 1A/1 P-10-12

AUTHORS: Bashkirov, A. N., Kagan, Yu. B., Zliger, G. A. 62-58-4-21/32

TITLE: Composition of Products Obtained by the Synthesis of  
Amines of Carbon-, Hydrogen- and Ammonia Monoxide  
(Sostav produktov sinteza aminov iz okisi ugleroda,  
vodoroda i amniaka)

PERIODICAL: Izvestiya Akademii Nauk SSSR, Otdelenie Khimicheskikh  
Nauk, 1958, Nr 4, pp. 504-506 (USSR)

ABSTRACT: Already earlier the authors realized the synthesis of  
the alkyl amines of CO and ammonia in the presence of  
molten iron catalysts. The synthesizing products ob=  
tained contained up to 25% aliphatic amines. Further=  
more a perfection of the used catalysts was carried  
out. A stable, active and rather selective catalyst  
was found. In the presence of the catalyst the authors  
synthesized on most favorable conditions: from 1 m<sup>3</sup>  
120,0 grams of synthesis products (without water).  
Of these were 54,0% alkylamines (30% of which in liquid  
and 21,0% in gas state). Tables 1-4 give information  
on the results of the elementary analysis of some

Card 1/2

The Composition of the Synthetization Products of the  
Amines of Carbon-, Hydrogen- and Ammonia Monoxide 52-58-4-21/32

fractions of the distillation and the physical con-  
stants of some fractions.

There are 4 tables and 12 references, 3 of which are  
Soviet.

ASSOCIATION: Institut nefti Akademii nauk SSSR (Petroleum  
Institute, AS USSR)

SUBMITTED: November 10, 1957

AVAILABLE: Library of Congress

1. Catalysts--Synthesis--Study and teaching

Card 2/2

AUTHORS: ~~Kagan, Yu. B., Bashkirov, A. N.,~~ SCV/62-55-10-19/25  
~~Kryukov, Yu. B., Loktev, S. M., Orlova, N. A.~~

TITLE: On the Mechanism of the Catalytic Efficiency of Fused Iron Catalysts in the Synthesis of CO and H<sub>2</sub> (O mehanizm kataliticheskogo deystviya plavlykh zheleznykh katalizatorov sinteza iz CO i H<sub>2</sub>)

PERIODICAL: Izvestiya Akademii nauk SSSR. Otdeleniye khimicheskikh nauk, 1953, Nr 10, pp 1274 - 1275 (USSR)

ABSTRACT: In an earlier paper the authors showed that immediately after the reduction (by hydrogen at 1000°) fused iron catalysts in the hydrocarbon synthesis of CO and H<sub>2</sub> are not active any more. Only under the working conditions of the synthesis when the gas mixture CO+H<sub>2</sub> is passed through the catalyst gradually becomes active (for 13-20 hours). This phenomenon may be explained by a number of simultaneous reactions competing with each other. Due to the course of these reactions competing with each other the metallic iron regenerates often (under the conditions of the synthesis) from its

Card 1/2

On the Mechanism of the Catalytic Efficiency of Fused Iron Catalysts in the Synthesis of CO and H<sub>2</sub> SCV/C2-53-10-19/25

compounds, and at the surface of the operating catalyst the dynamic equilibrium of the surface phases of different chemical structure is obtained. As a consequence of these processes the activation of the catalyst occurs. Neither the iron itself nor compounds that might be formed from it are the reason for the activation of the catalyst surface. The hypothesis formed for the chain mechanism of the catalytic efficiency of iron catalysts (according to which the synthesis of CO and H<sub>2</sub> is caused by the reactions of carbon and hydrogen monoxide with iron and its compounds on the surface of the operating catalyst) was described in detail by the authors. There are 1 table and 1 reference, which is Soviet.

ASSOCIATION: Institut nefti Akademii nauk SSSR (Petroleum Institute AS USSR)

SUBMITTED: April 8, 1958  
Card 2/2

KAGAN, Yu. B.

5(3) 21(8) IMAGE 1 BOOK EXPLORATION 07/2001

Academy of Sci. USSR. Institute petro-	
logy, t. 12 (Transactions of the Petroleum Institute, USSR. Academy of Sci. USSR), Vol. 12, Moscow, Izdat. Akad. Nauk SSSR, 1953, 359 p. Printed 11/52	
1,700 copies printed.	
Mr. S. S. Berezinov, Professor; Ed. of Published House: K. G.	
Reviewers' Tech. Ed.: V. V. Golikova.	
REVIEW: The book is intended for scientists, engineers, and technicians in the petroleum industry.	
SYNOPSIS: This collection of articles describes the results of studies on the chemistry and technology of petroleum and gas conducted in the Soviet Academy of Sciences Institute of Petroleum, Academy of Sciences, USSR, in 1956 and 1957. A new section "Petroleum Synthesis and Technology" has been included in the collection of articles. A list of publications published by the Institute in 1956 and 1957 and a list of dissertations of the Doctor's and Candidate's degrees presented in 1956 and 1957 at open sessions of the Academy of the USSR, the All-Union Petroleum Institute, Academy of Sciences, USSR, are given. In the Introduction, V. V. Korshakova, I. A. Mar'yav, and V. I. Shchukina, "Change in the Activity of Stille Gel in the Chromatographic Separation of Hydrocarbons."	
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12. D. P. [de] M. A. M. Bakhitov, and V. V. Smirnits. Determination of the Content of Primary and Secondary Higher Alcohols by the Dehydration Method	397
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5(2)

SPT/156-55-1-42/54

AUTHORS: Bashkirev, A. N., Kamzolkin, Ye. V., Kagan, Yu. B.

TITLE: On Catalysts of the Decomposition of Carbon Monoxide (O katalizatorakh reaktsiy razlozheniya okси углерода)

PERIODICAL: Nauchnyye doklady vyschey shkoly. Khimiya i khimicheskaya tekhnologiya, 1959, Nr 1, pp 162 - 165 (USSR)

ABSTRACT: For the systematic investigation of the influence of catalysts on the reactions: 1)  $2 \text{Fe} + 2 \text{CO} \rightarrow \text{Fe}_2\text{C} + \text{CO}_2$ , and 2)  $2\text{CO} \rightarrow \text{C} + \text{CO}_2$ , melted iron catalysts with activating additives ( $\text{Al}_2\text{O}_3$ ,  $\text{SiO}_2$ ,  $\text{K}_2\text{O}$ ,  $\text{Cr}_2\text{O}_3$ ,  $\text{B}_2\text{O}_3$ ,  $\text{MoO}_3$ ,  $\text{V}_2\text{O}_5$ ,  $\text{MgO}$ ,  $\text{MnO}$ , and combinations thereof), as employed in the synthesis  $\text{Co}+\text{H}$ , were investigated. The results are listed in a table. Carbide formation (reaction 1) and reaction 2 were accelerated by  $\text{Al}_2\text{O}_3$  and  $\text{V}_2\text{O}_5$ .  $\text{K}_2\text{O}$  accelerates only reaction 1.  $\text{MoO}_3$  is ineffective,  $\text{Cr}_2\text{O}_3$ ,  $\text{FeCr}$ , and in particular  $\text{SiO}_2$  and  $\text{B}_2\text{O}_3$  exercise inhibitory effects. With several additives to the catalyst, the properties have, as a rule, a cumulative effect.

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On Catalysts of the Reactions of the Decomposition of      SUV/156-59-1-42/54  
Carbon Monoxide

Only FeCr, which, when added exclusively, inhibits the reaction, increases the reaction velocity in the presence of other additives. Thus an iron catalyst with kaolin (as an  $\text{SiO}_2$  vehicle),  $\text{K}_2\text{O}$  and FeCr showed the highest reactivity.

In this case, even  $\text{B}_2\text{O}_3$  inhibits only reaction 2. A further test series concerned industrial iron catalysts, which were also given additives. The table of the results shows that also in this case the same rules apply. There are 2 tables and 1 Soviet reference.

ASSOCIATION: Kafedra neftekhimicheskogo sinteza i iskusstvennogo zhidkogo topliva Moskovskogo instituta tonkoy khimicheskoy tekhnologii im. M. V. Lomonosova (Chair of Petroleum-chemical Synthesis and Artificial Liquid Fuels of the Moscow Institute of Fine Chemical Technology imeni M. V. Lomonosova)

SUBMITTED: October 6, 1958

Card 2/2

SOV/62-59-7-34/38

5(3)  
AUTHORS: Kagan, Yu. B., Bashkirov, A. N., Klijer, G. A., Ferzikov,  
Yu. I.

TITLE: Transformation of n-Butyl Amine Under Synthesis Conditions  
From Carbon Oxide and Hydrogen (Prevrashcheniye n-butilamina  
v usloviyakh sinteza iz okisi ugleroda i vodoroda)

PERIODICAL: Izvestiya Akademii nauk SSSR. Otdeleniye khimicheskikh nauk,  
1959, Nr 7, pp 1345 - 1346 (USSR)

ABSTRACT: It had been found in previous investigations that when syn-  
thesizing n-butyl amine from CO, H<sub>2</sub> and NH<sub>3</sub> in the presence  
of iron catalysts, di- and trialkyls are formed in addition  
to monoalkyl amines. Moreover, the formation of the di- and  
trialkyls was found to be dependent in the course of reaction  
by the facilitated reacting possibility of primary amines with  
oxygen-containing products from the hydrogenation of the  
carbon oxide (alcohols, aldehydes, etc). The conclusion was  
drawn therefrom that the secondary and tertiary amines may be  
obtained from CO, H<sub>2</sub>, and the primary amine, here n-butyl  
amine. This conclusion is submitted to examination in the

Card 1/3

Transformation of n-Butyl Amine Under Synthesis  
Conditions From Carbon Oxide and Hydrogen

SUV/62-30-7-34/38

present paper. The synthesis conditions were the same as in references 1 and 2. A molten iron catalyst was also used in the investigation. The amines obtained were potentiometrically titrated. The following was investigated in the course of synthesis: the influence exerted by the amine addition rate upon the catalyst, by temperature, pressure, and the ratio CO/H in the initial mixture on the yield and the products of the synthesis. The data obtained are specified in the table. It may be observed from the latter that secondary and tertiary amines may be actually obtained in the manner described, and that, by changing the conditions, the reaction may be directed to the production of either secondary or tertiary amines. Increase in the concentration of n-butyl amine and temperature leads to the predominant formation of secondary amine, dilution of hydrogen, low temperature, and slow addition of n-butyl amine on the catalyst for the formation of tertiary amine. Rising pressure increases the formation of the two amines. At 80-150° the largest percentage yield of secondary and tertiary amines is obtained (45%). The additionally obtained primary amine (50%) is caused by disproportionation. There are 1

Card 2/3

Transformation of n-Butyl Amine Under Synthesis  
Conditions From Carbon Oxide and Hydrogen

S07/C2-50-7-31/30

Table and 10 references, 7 of which are Soviet.

ASSOCIATION: Institut nafti Akademii nauk SSSR (Institute of Petroleum  
of the Academy of Sciences, USSR)

SUBMITTED: January 19, 1959

Card 3/3

KAGAN, Yu.B.; BASHKIROV, A.N.; MOROZOV, N.G.; KRYUKOV, Yu.B.; ROZOVSKIY, A.Ya.

Hydrogenating capacity of fused iron catalysts in the synthesis  
from CO and H<sub>2</sub>. Trudy Inst.nefti 13:167-179 '59. (MIRA 13:12)  
(Catalysts) (Hydrogenation)

BASHKIROV, A.N.; LOKTEV, S.M.; KAGAN, Yu.B.; SABIROVA, G.V.

Hydrogenation of compounds containing a carbonyl group (over fused  
iron catalysts). Trudy Inst.nefti 13:180-195 '59. (MIRA 13:12)  
(Carbonyl compounds) (Hydrogenation)  
(Catalysts)

KAGAN, Yu. B.

2/195/00/001/002/006/010  
B002/005?

AUTHORS: Kostikov, Ya. B.; Babkinov, A. G.; Leshov, L. G.;  
Bartovits, V. F.; Stepanov, N. P.; ~~etc.~~

TITLE: Generation of Ionic Carboide Under the Conditions of the  
Synthesis of Hydrocarbons from Carbon Monoxide and Hydrogen

PUBLICATION: Elektro i metali, 1960, Vol. 17, No. 2, pp. 274 - 281

NOTE: The present paper was presented at the All-Union Conference on  
Inorganic Catalysis in December 1959. The authors attempted to explain the  
role played by carbides as intermediate compounds in the synthesis of  
hydrocarbons. They used a standard type catalyst with charcoal as  
a carrier, which was reduced at 1000°C and activated at 300°C and 20 atm  
with the addition of either  $\text{CO} + \text{H}_2$  (1 : 1), which contained  $\text{C}^{14}$ . The  
catalyst, saturated with radioactive iron carbide, was then treated with  
either  $\text{CO} + \text{H}_2$ . The radioactivity of the products formed was then measured.  
The authors found that nearly the following reactions took place in the

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2/195/00/001/002/006/010  
B002/005?

Characterization of Ionic Carbide Under the  
Conditions of the Synthesis of Hydrocarbons  
from Carbon Monoxide and Hydrogen

NOTE: Negotiations to submit, exchange of C isotopes between  
 $\text{CO}$  and carbide. The ratio of their isotopes varies as compared to that  
of the syntheses reaction, or 3000 CO molecules, only carbide 110  
molecules of 3000 CO molecules, only five are formed by carbide hydrogenation.  
Thus, only 0.01% of the hydrocarbons etc.  $\text{C}^{14}$  was formed under  
the action of carbide. Thus, due to what the syntheses associated to which  
carbide products are intermediates in hydrocarbons synthesis from  $\text{CO}$  and  
 $\text{H}_2$ . There are 2 figures, 2 tables, and 22 references 13 Soviet, 5 US,  
1 British, and 3 German.

DISSEMINATION: English translation available at USSR Institute of  
Petroleum Synthesis of the USSR (Institute of  
Petroleum Synthesis)

AUTHOR: January 25, 1960

Card 2/2

8/195/60/001/003/008/013  
B013/B058

AUTHORS: Kagan, Yu. B. Bashkirov, A. N., Kamzolkina, Ye. V.,  
Loktev, S. M.

TITLE: On the Activation Process of Molten Iron Catalysts for  
for CO and H<sub>2</sub> Synthesis Under the Effect of the Reaction  
Mixture

PERIODICAL: Kinetika i kataliz, 1960, Vol. 1, No. 3, pp. 393 - 400

TEXT: The activation of molten iron catalysts for the hydrocarbon  
synthesis from CO and H<sub>2</sub> under the effect of the reaction mixture was  
studied in this paper. The following catalysts were used:

- 1) 100Fe<sub>3</sub>O<sub>4</sub> + 6Al<sub>2</sub>O<sub>3</sub> + 4.2SiO<sub>2</sub> + 1.2K<sub>2</sub>O + 0.3Cr;
- 2) 100Fe<sub>3</sub>O<sub>4</sub> + 6Al<sub>2</sub>O<sub>3</sub> + 4.2SiO<sub>2</sub> + 1.2K<sub>2</sub>O + 0.5V;
- 3) 100Fe<sub>3</sub>O<sub>4</sub> + 6Al<sub>2</sub>O<sub>3</sub> + 4.2SiO<sub>2</sub> + 1.2K<sub>2</sub>O + 1.0B<sub>2</sub>O<sub>3</sub>. They were reduced

Card 1/4

On the Activation Process of Molten Iron      8/195/60/001/003/008/013  
Catalysts for CO and H<sub>2</sub> Synthesis Under the      B013/B058  
Effect of the Reaction Mixture

within 1.5 hrs in hydrogen current at 1000°C. The study was made in a highpressure apparatus (Ref. 4) in the laboratory. For each of the catalysts studied, the lowest temperatures and pressures were initially chosen, at which, over the freshly and reduced catalysts (in comparable time intervals), a high degree of transformation of the carbon monoxide (84 to 86%) entering at a volume rate of the initial gas (CO and H<sub>2</sub> 1:1) of  $\sim 1500 \text{ h}^{-1}$  was obtained. The catalysts were gradually activated under these conditions. The duration of the tests varied. The tests of catalysts of equal composition were conducted under the same conditions and in the same reaction vessel. The indices of the synthesis were well reproducible. The results determined could therefore also be compared with each other. The samples were hydrogenated after termination of the synthesis test. Subsequently, the hydrogenated samples were treated with CO at atmospheric pressure, a volume rate of 800  $\text{h}^{-1}$  and temperatures by 10° higher than at the end of the synthesis test, with carbide being formed. The studies

Card 2/4

On the Activation Process of Molten Iron  
Catalysts for CO and H<sub>2</sub>Synthesis Under the  
Effect of the Reaction Mixture

S/195/60/001/003/008/013  
B013/B058

the activation of the catalyst for the synthesis, its treatment at the synthesis temperature is suited best. In this case the catalysts get specially active, but simultaneously show a lower activity with regard to CO decomposition. There are 3 figures, 4 tables, and 6 Soviet references.

ASSOCIATION: Institut neftekhimicheskogo sinteza AN SSSR  
(Institute of Petrochemical Synthesis AS USSR)

SUBMITTED: January 19, 1960

Card 4/4

KAGAN, Yu.B.; BASHKIROV, A.N.; KLIGER, G.A.

Amination of n-butanol over iron catalysts under hydrogen pressure.  
Izv.AN SSSR Otd.khim.nauk no.3:468-473 Mr "61. (MIRA 14:4)

1. Institut neftekhimicheskogo sinteza AN SSSR.  
(Butyl alcohol) (Amination)

KAGAN, Yu.B.; BASHKIROV, A.N.; KLIGER, G.A.; CHZHOU CHZHAO-DI [Chou Chao-ti];  
MAK, N.Ye.

Reaction between octyl alcohols and ammonia under the hydro-  
gen pressure on a fused iron catalyst. Neftekhimia 1 no.3:  
403-410 My-Je '61. (MIRA 16:11)

1. Institut neftekhimicheskogo sinteza AN SSSR i Institut  
tonkoy khimicheskoy tekhnologii imeni Lomonosova.

KAGAN, Yu.B.; BASHKIROV, A.N.; KLIGGER, G.A.; CHZHOU CHZHAO-DI  
[Chou Chao-ti]; MAK, N.Ye.

Effect of the molecular weight of alcohol on the process  
of its amination. Neftekhimika 1 no.4:555-563 Jl-Ag '61.  
(MIRA 16:11)

1. Institut neftekhimicheskogo sinteza AN SSSR i Moskovskiy  
institut tonkoy khimicheskoy tekhnologii imeni Lomonosova.

KAGAN, Yu.B.; ROZOVSKIY A. Ya.; KRYUKOV, Yu.B.

Mechanism of the action of fused iron catalysts in the synthesis of  
organic compounds from CO and H<sub>2</sub>. Kin. i kat. 2 no.1:55-60 Ja-F '61.  
(MIRA 14:3)

1. Institut neftekhimicheskogo sinteza AN SSSR.  
(Carbon monoxide) (Hydrogen) (Catalysis) (Iron)

ROZOVSKIY, A.Ya.; BASHKIROV, A.N.; KAGAN, Yu.B.; POKROVSKAYA, Ye.G.

Water and water vapor oxidation of the iron catalysts for  
synthesis from CO and H<sub>2</sub>. Kin.i kat. 2 no.6:830-837 N-D '61.  
(NIRA 14:12)

1. Institut neftekhimicheskogo sinteza AN SSSR.  
(Carbon monoxide)  
(Hydrogen) (Catalysts, Iron)

KAGAN, Yu.B.; BASHIKIROV, A.N.; KLIGER, G.A.; ROZOVSKIY, A.Ya.

Certain problems involved in the microkinetics of amination of  
alcohols on fused iron catalysts. Neftekhimiia 2 no.2:253-256  
Mr-Ap '62. (MIRA 15:6)

1. Institut neftekhimicheskogo sinteza AN SSSR.  
(Alcohols) (Amination)

KLIGER, G.A.; BASHKIROV, A.N. LYUY GUAN-YUY [Li Kuang-yü]; LESIK, O.A.;  
SEZINGER, N.N.; KAGAN, Yu.B.

Method of analyzing products of reaction between aliphatic  
alcohols and alkyl amines. Neftekhimia 2 no.1:121-126 Ja-F  
'62. (MIRA 15:5)

1. Institut neftekhimicheskogo sinteza AN SSSR.  
(Alcohols) (Amines)

KLIGER, G.A.; BASHKIROV, A.N.; LYUY GUAN-YUY [Lü Kuang-yü]; KAGAN, Yu.B.

Effect of the structure and molecular weight of initial compounds  
on the interaction of aliphatic alcohols with primary alkyl  
amines. Neftekhimiia 2 no.3:384-390 My-Je '62. (MiRA 15:8)

1. Institut neftekhimicheskogo sinteza AN SSSR.  
(Alcohols) (Amines)

KAGAN, Yu.B.; BASHKIROV, A.N.; KLIGER, G.A.; LYUY GUAN-YUY [Lü Kuang-yü]

Synthesis of secondary alkyl amines from aliphatic alcohols  
and primary amines. Neftekhimiia 2 no.3:391-397 My-Je '62.  
(MIRA 15:3)

1. Institut neftekhimicheskogo sintesa AN SSSR.  
(Amines) (Alcohols)

ROZOVSKIY, A.Ya.; BIRYUKOVICH, M.M.; IVANOV, A.A.; LIBEROV, L.G.;  
BUTYUGIN, V.K.; KAGAN, Yu.B.; KRYUKOV, Yu.B.; BASHKIROV, A.N.

Mechanism of the carbide-forming reaction of fused iron  
catalysts for synthesis from CO and H<sub>2</sub>. Neftekhimiia.  
3 no.1:97-103 Ja-F '63. (MIRA 16:2)  
(Iron catalysts) (Iron carbides)  
(Chemistry, Organic—Synthesis)

ROZOVSKIY, A.Ya.; IVANOV, A.A.; KAGAN, Yu.B.; BASHKIROV, A.N.

Kinetics of reactions involving the solid phase. Part 2: Hydrogenation  
of iron carbides. Kin.i kat. 4 no.1:97-108 Ja-F '63. (MIRA 16:3)

1. Institut neftekhimicheskogo sinteza AN SSSR i Institut tonkoy  
khimicheskoy tekhnologii imeni M.V.Lomonosova.  
(Iron carbides) (Hydrogenation)

ROZOVSKIY, A.Ya.; BIRYUKOVICH, M.M.; IVANOV, A.A.; KAGAN, Yu.B.;  
BASHKIROV, A.N.

Kinetics of reactions involving the solid phase. Part 3:  
Carbide formation in fused iron catalysts induced by carbon  
monoxide. Kin. i kat. 4 no.3:373-381 My-Je '63.  
(MIRA 16:7)

1. Institut neftekhimicheskogo sinteza AN SSSR.  
(Carbides) (Iron catalysts)  
(Chemical reaction, Rate of)

KLIGER, G.A.; BASHKIROV, A.N.; BEZINGER, N.N.; KAGAN, Yu.B.

Method for analyzing products obtained by the interaction  
of aliphatic alcohols with ammonia in the presence of  
hydrogen. Neftekhimia 1 no.3:397-402 My-Je '61.  
(MIRA 16:11)

1. Institut neftekhimicheskogo sinteza AN SSSR.

LOKTEV, A.M.; KAGAN, Yu.B.

Selective catalytic hydrogenation of aliphatic oxygen-containing compounds under high pressures. Neftekhimika 3 no.6:892-899 N-D '63. (MIRA 17:3)

1. Institut neftekhimicheskogo sinteza AN SSSR im. A.V.Topchiyeva i Novomoskovskiy khimicheskiy kombinat.

PONOMARENKO, A.T.; KAGAN, Yu.B.; KAMZOLKIN, V.V.

Device for measuring gas consumption under high pressure. Khim.  
i tekhn.topl. i masel 9 no.2:48-50 F '64. (MIRA 17:4)

1. Institut neftekhimicheskogo sinteza AN SSSR.

ACCESSION NR: AP4024408

8/0204/64/004/001/0106/0110

AUTHOR: Kagan, Yu. B.; Bashkirov, A. N.; Kliger, G. A.; Iu, Kuang-yu  
TITLE: Synthesis of secondary alkylamines from carbon monoxide, hydrogen and methylamine.

SOURCE: Neftekhimiya, v. 4, no. 1, 1964, 106-110

TOPIC TAGS: secondary alkylamine, synthesis, amination, process condition, reaction mechanism, methylalkylamine

ABSTRACT: Continuing earlier work (Yu. B. Kagan, A. N. Bashkirov, G. A. Kliger, Yu. I. Yermakov. Izv. AN SSSR, Otd. khim. n. (1959), 1345), the interaction of CO, H<sub>2</sub> and methylamine to form secondary alkylamines was studied. Using a fused iron catalyst, the optimum synthesis conditions are 100-140 atmospheres pressure, 1250 hours<sup>-1</sup>, volume flow of the initial gas mixture, a temperature of 168-175 C, and an H<sub>2</sub>:CO:CH<sub>3</sub>NH<sub>2</sub> ratio of 4:1:1.25. The yield of secondary amines was 77-78.5 mol.%. The product also contained small amounts of alcohols and carbonyl com-

ounds, indicating the synthesis probably proceeded in two stages; (1) the formation of the oxygen-containing compounds from CO and H<sub>2</sub>, (2) followed by

Card 1/2

ROZOVSKIY, A. Ya.; BIRYUKOVICH, M.M.; IVANOV, A.A.; KAGAN, Yu.B.; BASHKIROV,  
A.N.

Kinetics and mechanism of the carbidizing treatment of iron  
catalysts for synthesis from CO and H<sub>2</sub>. Neftekhimiia 4 no.2:  
269-274 Mr-Ap'64 (MIRA 17:8)

1. Institut neftekhimicheskogo sinteza AN SSSR imeni Topchiyeva.

VULAKH, Yu.L.; LOKTEV, S.M.; KAGAN, Yu.B.

Esterification of aliphatic alcohols with sulfuric acid. Neftekhimika  
4 no.5:780-788 S-0 '64.  
(MIRA 18:1)

1. Institut neftekhimicheskogo sinteza imeni A.V.Topchiyeva AN SSSR  
i Novomoskovskiy khimicheskly kombinat.

KAGAN, Yu.B.; ROZOVSKIY, A.Ya.; SLIN'KO, M.G.; PONOMARENKO, A.T.

Kinetics of heterogeneous catalytic reactions as a function of  
ignition conditions. Part 2: Reaction of an arbitrary order.  
Kin. i kat. 5 no.6:1111-1114 N-D '64.

(MIRA 18:3)

1. Institut neftekhimicheskogo sinteza imeni Topchlyeva AN SSSR  
i Institut kataliza Sibirskogo otdeleniya AN SSSR.

KAGAN, Yu.B.; ROZOVSKIY, A.Ya.; SLIN'KO, M.G.; PONOMARENKO, A.T.

Study of the kinetics of catalytic reactions based on the  
conditions of ignition. Part 1: Zero-order reactions. Kin.  
i kat. 5 no.5:920-926 S.O '64. (MIRA 17:12)

1. Institut neftekhimicheskogo sinteza imeni Topchiyeva AN SSSR  
1 Institut kataliza Sibirskego otdeleniya AN SSSR.

KAMZOLKIN, V.V.; KRYUKOV, Yu.B.; KAGAN, Yu.B.

Prospective trend of the petroleum chemistry. Vest. AN SSSR 34  
no.11:60-65 N '64. (MIRA 17:12)

1. Institut neftekhimicheskogo sinteza im. A.V. Topchiiyeva AN SSSR.

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KAGAN, S.Z.; KOVALEV, Iu.N.; KACAN, Yu.B.; OPOLOVA, N.A.

Studying the extraction of higher alcohols from their mixtures  
with hydrocarbons. Trudy MFTI no. 40. 128-133 - 1959.  
(MIRA 18:12)

KUTEPOVA, A.I.; GRISHKO, N.I.; KAGAN, Yu.B.; LOKTEV, S.M.; MAL'TSEVA, R.P.;  
SHTEKKER, O.A.

Preparation of phthalate plasticizers on the base of the wide  
fractions of C<sub>5</sub>-C<sub>12</sub> alcohols. Plast. massy. no.10:22-24 '65.  
(MIRA 18:10)

KAGAN, Yu.L. (Moskva, B-108, Staromonetnyy per., d.22, kv.3); BAKULEVA, L.P.,  
kand.med.nauk

Possibility of studying the contractile activity of the uterus and  
Fallopian tubes by roentgenokymography. Vest.rent.i rad. 34 no.5:  
52-55 S-0 '59. (MIRA 13:3)

1. Iz kafedry rentgenologii i radiologii (zav. - prof. V.A. D'yachenko)  
i kafedry akusherstva i ginekologii (zav. - prof. I.Y. Zhordania) II  
Moskovskogo meditsinskogo instituta imeni N.I. Pirogova.  
(UTERUS radiography)  
(FALLOPIAN TUBES radiography)

KHOMYAKOV, Yu.S. (Moskva, v-95, 2-y Kazachiy pereulik, d.3, kv.2);  
KAGAN, Yu.L.

X-ray diagnosis of gastric burns. Vest.rent.1 rad. 35 no.1:  
3-6 Ja-# '60. (MIRA 13:6)

1. Iz kafedry rentgenologii i radiologii (zav. - prof. V.A.  
D'yachenko) II Moskovskogo meditsinskogo instituta imeni N.I.  
Pirogova.

(CAUSTICS)  
(STOMACH wds. & inj.)

SADOVSKAYA, I.A.; KAGAN, Yu.L.

Cleidocranial dysostosis diagnosed in a newborn infant. Pediatrichia  
38 no.10:75-77 0 '60. (MIRA 13:11)

1. Iz rodil'nogo doma No.7 imeni Grauermana (glavnnyy vrach  
Ye.I. Bulanova). (DYSOSTOSIS)

DEMIDKIN, P.N.; KAGAN, Yu.L.

First experience in using the new Soviet intensifying screens for  
roentgenography in obstetrical practice. Vop. okh. mat. i det. 7 no.3:  
62-63 Mr '62. (MIRA 15:5)

1. Iz rentgenovskogo otdeleniya (zav. - kand.med.nauk P.N.Demidkin)  
Moskovskogo oblastnogo nauchno-issledovatel'skogo instituta akusherstva  
i ginekologii (dir. - zasluzhennyj vrach RSFSR O.D.Matspanova, nauchnyj  
rukovoditel' - prof. A.V.Lankovits).  
(RADIOGRAPHY--EQUIPMENT AND SUPPLIES)  
(OBSTETRICS)

C A

3

LUMINESCENCE AND FLUORESCENCE  
Luminescence of ions in the positive column of a glow discharge. Yu. M. Kagan and S. R. Prish. *J. Russ. Phys. Chem. Soc.* 11, 260-9 (1911).—The luminescence of A in a silent discharge current densities up to 65 amp./sq. cm. is illustrated for 7 lines of A I and 20 of A II by 4 figs. The intensity of the A II lines from 4000-4000 Å. is proportional to the square of the d. of the discharge current. The discharge is considered as at the nonequilibrium limit relative to the excitation of ions. The intensity of the A I lines rapidly reaches satn. at around 20-25 amp./sq. cm. P. H. Rathmann

Physics Inst., Hungarian State U.

ASSISTANT CHIEF LIBRARIAN CLASSIFICATION

KAGAN, Yu. M.

"Spectroscopic Investigation of Ions in the Positive Column of a Glow Discharge," Zhur. eksper. i teoret. fiz., 12, No.9, 1942

Leningrad State U.

c A

Spectroscopic investigation of ions in the positive column of a glow discharge. S. E. Frish and Yu. M. Kagan. J. Phys. (U. S. S. R.) 7, 208 (1939) (in English).—Observations on the luminescence of a positive column of a glow discharge in A and Ne at large c. d. disclosed the appearance of spark lines along with the arc lines. A quadratic dependence was found for the intensity of the A II and Ne II lines on the strength of the discharge current. The dependence of the intensity of the A II lines on the pressure at a const. c. d. was also studied. By a study of the shape of the A I and A II lines it was established that the energy of the random motion of the ions in the discharge is of the order of 0.3 e. v., which is larger than the thermal energy of neutral atoms. With increase of the strength of the discharge current the difference between the energy of the atoms and that of the ions is diminished. The velocity distribution of the ions is markedly asymmetrical, the direction of the elec. field being the axis of symmetry. The drift velocity of the ions is comparable with the velocity of their random motion.  
F. B. Rathmeyer

Physics Inst., Ferganid State U.

6602

24.2/20  
 AUTHORS: Granovskiy, V.I., Luk'yano, S.O., Sirotnikov, G.V. and Sirotnikova, T.G.  
 TITLE: Report on the Second All-Union Conference on Gas Discharges

KAGAN, Yu.M.

PERIODICAL: Radiotekhnika i elektronika, 1959, Vol. 4, Nr. 8,  
 pp 1339 - 1350 (USSR)  
 ABSTRACT: The conference was organized by the Ac.Sc.USSR, the Ministry of Higher Education and Moscow State University. It was opened by the chairman of the organizing committee, N.A. Lachinovich. Academician. During the plenary sessions of the conference, a number of survey papers were delivered. L.A. Aksel'movich read a paper on "Production of Ultra-high Temperatures in Plasma". A survey of the optical method of measurement was given in the papers by V.A. Fabrikant and S.Z. Prian. S. Brown of the Massachusetts Institute of Technology gave a survey of the high-frequency methods of the investigation of stationary and non-stationary plasma (see p. 1244 in this issue of the journal).

N.V. Fedorenko read a paper entitled "Ionization and Ionization-Neutralization During Atomic Processes". X

L.A. Aksel'movich and Yu.N. Kachalov, with "Elementary Processes of Detonation", the notion of zones of the role of resonance-recharging in the kinetics of zones". A paper by Ye. Bedroev (Bulgaria) dealt with the development of sparks (corona-leader, main channel and the final channel).

B.F. Myrfield gave a survey of the ignition processes of the discharges in highly rarified gases. The mechanism of the breakdown of a high-vacuum gap was elucidated in a paper by V.L. Granovskiy.

I. Tondre (USA) expounded a theory of the motion of electrons in a magnetic trap (see p. 1316 of this journal). A. Kedem (Israel) described a number of experiments on non-stationary plasma conducted by himself.

M. Schubert (Western Germany) gave a generalized theory of plasma. The conference was divided into six sections of which one section was presided over by L.A. Sere and was concerned with the elementary processes in ion discharges. The following papers were read in this section:

Ish. M. Projai - "Transformation of Positive Ions Rate, Negative Ones in Rarified Gases".

Ye. M. Fesel, with V.A. Andrunin and D.V. Pilishevskiy - "Capture and Loss of Electrons During the Collision of Fast Atoms or Carbon and Hydrogen with the Molecules of Gases".

M.V. Fadchenko et al. - "Dissociation or Molecular Ions of Hydrogen During Collisions in Gase".

I.P. Flaks and Ya.K. Kholmyan - "Capture Cross-sections of Electrons in Multicharge Ions in a Rarefied Gase".

R.H. Kuhnlich et al. - "Experimental Investigation of the Measuring Scattering in Certain Single-system Gases and Metal Vacuums".

O.B. Piriey - "Qualitative Investigation of Inelastic Collisions of Atoms".

L.H. Wohlberg - "Infective Excitation Cross-sections of the Artificial Ions at Potassium and Arsenic".

Card 15 I.P. Zhdanovskiy and S.M. Kishkin - "Some Results of the Investigation of the Optical Functions of the Emission Bands of a Negative System".

A.A. Vorob'ev and A.G. Vinogradov - "Investigation of the Scattering of the Electrons in a Betatron Chamber".

The second section was presided over by B.M. Klyarov and was devoted to the problems of the electrical breakdown in rarified gases and in high vacuum. The following papers were read in this section:

Q.Ye. Makar-Shanov and Yu.A. Matlakovsky - "Dielectric Control of the Ignition of Glow-discharge Tubes" (see

p. 1274 of the journal).

S.Y. Pletnev et al. were concerned with the breakdown in a high-voltage secondary rectifier (see p. 1275 of the journal).

I.G. Guseva - "Emission of the Discharge in Non-uniform

Fields at Low Gas Pressures" (see p. 1260 of the journal).

A.S. Dobrovolskaya and B.N. Il'yashenko - "The Dielectric Properties of Solid and a Flame at the Pressure of  $10^{-3} - 1 \text{ mm Hg}$ ".

F  
G

2011. SPECTROSCOPIC STUDY OF ELECTRIC CRACKING OF METHANE.  
Frisch, S. E. and Kagan, Y. M. (Bull. Acad. Sci. U.S.S.R., Ser. Phys., 1945, 2, 238). The emission spectrum of CH<sub>4</sub> in an elec. discharge, at 35 mm. pressure, shows mol. bands C<sub>2</sub>, CH, and CN (impurities). In the visible part one finds a continuous spectrum and the lines H<sub>β</sub> and H<sub>α</sub>. Stopping the flow of methane resulted in enhanced hydrogen spectrum; the CH bands remained unchanged, CN weakened. With both the gas flowing and at rest, the CH bands show many rotation lines, indicating high rotational energies of the CH mol. From the continuous spectrum, the temp. was detd. to be 1200° K. It can be concluded that CH mols. are present as intermediate products and that the process deviates from equilibrium with respect to temperature.

C. A.

ASB-SLA METALLURGICAL LITERATURE CLASSIFICATION

C 4

- Polytechnic Inst., Leningrad, U.

Spectroscopic study of the motion of ions in the plasma  
S. E. Frish and Yu. M. Kagan (Leningrad Univ.). Zhur.  
*Exptl. Teoret. Fiz.*, 17, 577-81 (1947).—Interferometric  
measurements made on spectral lines of A II along a  
quartz capillary in which the discharge takes place, show  
a displacement absent on lines of A I. This displacement  
of the magnitude of  $0.001 \pm 0.001$  Å is attributed to the  
Doppler effect caused by the translatory movement of ions  
along the capillary with the velocity of  $1 \times 10^8 - 2.3 \times$   
 $10^8$  cm./sec. This is confirmed by: (a) the ratio of dis-  
placements of lines 5002 and 4230 is equal to the ratio of  
wave lengths as predicted by the theory; (b) the displace-  
ment is proportional to the cosine of the viewing angle with  
the axis of the capillary; (c) the velocities calcd. from the  
displacements at different pressures (0.2-3.5 mm.) and  
discharge currents (100-400 ma) have been compared  
with a theoretically derived formula,  $(\Delta v)^2 = 7 \times$   
 $10^9 T/27 D^2 [E/p(1 + (E_r/E))]^{1/2}$  ( $E$  measured tube  
gradient,  $E_r$  calcd. radial gradient) and found to be in satis-  
factory agreement. S. Pakser

KAGAN, YU. M. Doc Phys-Math Sci -- (diss) "On the motion of  
ions in plasma." Len, 1957. 15 pp 20 cm. (Min of Higher Edu-  
cation USSR. Moscow Order of Lenin and Labor Red Banner State  
Univ im M.V. Lomonosov), 100 copies  
(KL, 21-57, 97)

-3-

KAGAN, YU. M., DOCENT

Jan 18

USSR /Physics  
Discharges, Electric  
Spectrum Analysis

"Spectroscopy of Gas Discharge," S. Z. Frish, Corp

Mem. Acad Sci USSR, Docent Yu. M. Kagan, 29 pp

West Leningrad U<sup>n</sup> № 1

Theoretical survey of subject: (1) phenomena occurring in a gas discharge; (2) step-by-step activation of ions on positive pole of a gas discharge; (3) effect of spectroscopic study of movement of ions in plumes on positive pole; (4) effect of cascade migrations on form of activation functions; (5) role of second order impacts on luminescence of the lines during gas discharge; (6) application of gas discharge spectrum analysis of gas mixtures.

4/49T69

KAGAN, YU. M.

Jul/Aug 48

USSR/Physics

Ions  
Gas

"Movement and Form of Lines of Ions at the Positive  
Poles During Gas Discharge," S. E. Frish, Yu. M. Kagan,  
Sci Res Phys Inst, Leningrad State U, 4 pp

"Iz Ak Nauk SSSR, Ser Fiz" Vol XII, No 4  
[17-158-6]

[17-158-6]  
Introduces curves showing translational speeds of  
positive Ar, Kr, and Xe ions versus gas pressure for  
a single current strength of 400 ma. At three curves  
have approximately the same path: When pressure drops,  
speed initially increases and then reaches a maximum

53/49r88

Jul/Aug 48

USSR/Physics

(Contd)

and drops for lower pressures. Theoretical explanation  
of this phenomenon.

53/49r88

*Cy*

## PROCESSES AND PROCEDURES (CONT)

The influence of cascade transitions on the form of the excitation function of mercury lines. Yu. M. Kagan and V. M. Zakharova (Leningrad State Univ.) ZH. fiz. chisl. i Teorii. Fiz. (J. Exptl. Theoret. Phys.) 18, 537 (1948).—Two explanations have been proposed for the existence of two max. in the excitation functions of the  $6^3P_1 - 6^1S_0$  series of Hg lines. K. and Z. disprove Schaffernicht's suggestion (C.I. 24, 3021) that the hyperfine structure sublevels of  $6^1S_0$  may have different excitation functions by showing experimentally that the relative intensities of the hyperfine structure components

of Hg 4046 and Hg 4077 are not a function of the temp. of the Hg vapor. If the shape of the excitation function is due to the existence of a cascade transition from a higher level, it is necessary to explain why such transitions do not occur to other term levels. K. and Z. believe that this is due to the unusually low cross section of the  $6^1S_0$  level. Their exptl. dets. of  $Q_{ext}(V)$ , the effective cross section of the  $6^1S_0$  level indicated at the first max. in the excitation curve, gave the following values:  $7^1S_0$ : 0.015-0.020;  $8^1S_0$ : 0.006-0.004;  $9^1S_0$ : 0.010-0.013. C. F.

*Physics Inst.*

## ABR-SLA METALLURGICAL LITERATURE CLASSIFICATION

EIGHT DIVISIONS: 1. GENERAL INFORMATION 2. MATERIALS 3. PROCESSING 4. EQUIPMENT 5. CONSTRUCTION 6. DESIGN 7. OPERATIONS 8. COSTS

KAGAN, YU. M.

(3)

Spectroscopic study of the movement of ions in a plasma.  
II. S. K. Frish and Yu. M. Kagan [Leningrad State Univ.].  
*Zhur. Ekspd. i. Teor. fiz.* 18, 519-24 (1948); *cf. C.A.* 44,  
6200d.—By using the calibrating interferometer device of  
Fabry and Perot, F. and K. studied the contours of the  
spectral lines of ions and of neutral atoms of argon as a  
function of the strength of the discharge current and the  
pressure of the gas. Lines which are due to ions are much  
wider than are lines due to neutral atoms. Lines of ions are  
wider when viewed across the capillary than when viewed  
along the capillary. Using the half-widths of the lines, F.  
and K. calc'd. the ionic temps., and compared these with the  
temp. of the st. gas. During observations along the capil-  
lary, the course of the ionic temp. as a function of the pres-  
sure is analogous to that already previously established for  
the excited lines of ions. The results obtained are inter-  
preted from the point of view of transference of ions  
in an elec. field in a plasma. (7)  
Franz H. Rothmann

KAGAN, YU. M.

USSR/Physics - Gaseous Discharge

Nov/Dec 50

"Concentration of Excited Atoms in a Mercury Discharge," Yu. M. Kagan, N. P. Fenkin, Sci. Res Phys Inst, Leningrad State U imeni Zhdanov

"Iz Ak Nauk SSSR, Ser Fiz" Vol XIV, No 5,  
pp 721-6

Studies following dependences: electron temperature vs pressure; electron concentration vs pressure, discharge current strength; population of various electron levels vs pressure and electron concentration. Authors thank S. E. Frish and Ye. I. Pokrovskiy for their assistance.

170191

KAGAN, YU.

PA 1971102

USSR/Nuclear Physics - Atoms, Excitation Oct 51  
of

"Excitation of Atoms in Mercury Discharge," Yu.  
Kagan, N. Perkin

"Zhur Eksper i Teoret Fiz" Vol XXI, No 10, pp  
1182, 1183

Authors answer criticism of their article ("Iz  
Ak Nauk SSSR, Ser Fiz" 14, 1950) by Fabrikant and  
Yavorskiy (ibid. pp 1180, 1181) and emphasize  
that their theoretical curve of atomic energy is  
correct, despite objections by critics. They also  
deny having based their research on data by Fabri-  
kant and Yavorskiy. Submitted 30 Jun 51.

LC

1971102

PA 175T14

USSR/Electronics, Discharge in Gases 11 Jan 51

"Method of Investigation of Function of Distribution of Electrons Depending on Velocities in a Gas Discharge," Yu. M. Kagan, V. L. Fedorov, G. M. Malyshev, L. A. Cavallas

"Dok Ak Nauk SSSR" Vol LXXVI, No 2, pp 215-217

Describes method where curve of 2d deriv, necessary for computation, is obtained directly on oscillograph. Curves proved that the distribution function deviates from Maxwell's function near the cathode.

175T14

KAGAN, YU. M.

USR/Electricity - Discharge Tube

Apr 52

"Excitation of Ions in Mercury Discharge at Low Pressure," Yu. M. Kagan, V. M. Zakhareva, Lenin-grad State U-

"Zhur Elektr i Teoret Fiz" Vol XXXI, No 4, pp 400-405

Excitation of ions and atoms of mercury in the column of discharge at low pressure is studied. Spectroscopic observations confirm the assumption of Maxwell distribution of fast electrons according to velocities. The excitation of ions occurs as result of direct impact of atom and electron.

21ST22

Expts with shift of spectral lines lead to the conclusion that from the point of their origin the ions fall freely on the tube walls. Indebted to Prof S. E. Frish. Received 18 Jul 51.

21ST22

*RECORDED 7/17/77*

**USSR .**

Distribution of electrons according to speeds in the electrical discharge of mercury vapors. Yu. M. Kugan and G. M. Malyshev. Zhur. Tekh. Fiz. 23, 846 (1933).--  
A report of studies of the form of the distribution function for electron speeds in the mercury vapor discharge at different currents and pressures, at different distances from the cathode.

Gladys S. May

"APPROVED FOR RELEASE: 08/10/2001

CIA-RDP86-00513R000619910019-5

APPROVED FOR RELEASE: 08/10/2001

CIA-RDP86-00513R000619910019-5"

KAGAN, Yu. M.

USSR/Physics - Ion Current

21 Aug 53

"Theory of Ion Flow into a Probe at Low Voltages,"  
Yu. M. Kagan and V. I. Perel', Karelo-Finnish State  
Univ

DAN SSSR, Vol 91, No 6, pp 1321-1324

State that Langmuir's theory of probe characteristics (which proceeds from the assumption that a space-charge layer surrounds the probe and from the assumption that the plasma outside this layer is quasineutral and that the electrical field

275T96

equals zero and the motion of the particles is without order) leads to considerable divergences from experience when the theory is applied to the ion saturation flow toward a negatively charged probe. Acknowledge assistance of P. Ripatti and V. Slyasskiy in the measurements. Presented by Acad A. A. Lebedev 26 Jun 53.

## USSR/ Physics - Electron distribution

Card 1/1 Pub. 43 - 16/97

Authors : Zakharovz, V. M., and Kagan, Yu. M.

Title : Spectroscopic investigation of electron distribution according to the speeds  
in a positive gas discharge column

Periodical : Izv. AN SSSR. Ser. fiz. 18/2, page 254, Mar-Apr 1956

Abstract : The electron distribution was investigated in a positive discharge column  
in the presence of sodium - helium vapors at a discharge current intensity  
of 10 - 90 ma. It is assumed that the spectroscopic method will make it  
possible to obtain more accurate data on the speed distribution of fast elec-  
trons provided the discharge conditions are such that the secondary pro-  
cesses can be disregarded. The difference between the speed of normal  
electron distribution and the Maxwell distribution is briefly explained. One  
USSR reference (1941).

Institution : The A. A. Zhdanov State University, Physics Institute, Leningrad

Submitted : .....

KAGAN, Yu.M.; PERML', V.I.

Theory of the Langmuir ball-probe in the plasma. Dokl.AN SSSR  
(MLRA 7:3)  
95 no.4:765-768 Ap '54.

1. Karelo-finskiy gosudarstvennyy universitet. 2. Karelo-finskiy  
pedagogicheskiy institut. (Nuclear physics)

KAGAN, Yu. M.

USSR/Physics

Card 1/1 : Pub. 22 - 17/49

Authors : Kagan, Yu. M., and Perel', V. I.

Title : On the movement of positive ions in their own gas

Periodical : Dok. AN SSSR 98/4, 575-578, Oct. 1, 1954

Abstract : Computations of the velocities of positive ions moving in their gas in a strong and a weak-field are presented. Nine references (1946-1953). Graphs.

Institution : Karelo-Finnish State University, Karelo-Finnish Pedagogical Institute

Presented by : Academician A. A. Lebedev, February 25, 1954

KAGAN, Yu. M.

USSR/ Physics

Card 1/1 Pub. 177 - 7/13

Authors : Zokhoreva, V. M., and Kagan, Yu. M.  
Title : Line intensity distribution in secondary series in a positive sodium discharge column  
Periodical : Vest. Len. un. Ser. mat. fiz. khim. 10/2, 115-130, Feb 1955  
Abstract : The radiation of sodium diffusion series was investigated with respect to the change in discharge parameters to determine the deviations from the Maxwell electron distribution. The dependence of the electron temperature and electron concentration upon pressure and discharge current intensity is explained. The results obtained by measuring the line intensity distribution within secondary series at various pressures and current intensities are tabulated. The harmful effect of the electrophoresis phenomena, which exists during the discharge in a vapor-inert gas mixture, is discussed. Twenty references: 12 USSR, 3 German, 4 English and 1 Dutch (1930-1954). Graphs.

Institution : .....

Submitted : June 21, 1954

FD-2338

USSR/Physics - Probe theory

Card 1/1      Pub. 146 - 25/26

Author : Kagan, Yu. M.; Perel', V. I.

Title : Langmuir theory of probes (sondes)

Periodical : Zhur. eksp. i teor. fiz., 29, August 1955, 261-263

Abstract : Certain assumptions of the Langmuir theory of probes have been criticized earlier by the writers (DAN SSSR, 91, 1321, 1953) and others (R. Boyd, Proc. Roy. Soc., A 201, 329, 1950; F. Wenzl, Zs. angew. Phys., 2, 59, 1950). The task of the writers in the present note is to clarify the interrelationship between the Langmuir theory and the strict theory of the spherical probe proposed by the writers (DAN SSSR, 95, 765, 1954) for low pressures and for the case of negative probe potentials. They restate the principal assumptions under these conditions as given by V. L. Granovskiy (Elektricheskii tok v gaze [Electrical current in a gas], GIITL, 1952). Five references: e.g. Mott-Smith and Langmuir, Phys. Rev., 28, 1926.

Institution : Karelo-Finn State University

Submitted : April 9, 1955

FD-3271

USSR/Physics - Recharging of slow ions

Card 1/1 Pub. 146 - 30/44

Author : Kagan, Yu. M.; Perel', V. I.

Title : Cross section of overcharging of slow ions in its own gas

Periodical : Zhur. eksp. i teor. fiz., 29, No 6(12), Dec 1955, 884-886

Abstract : The authors compare the various values of overcharge of He, Ne, A, Kr, and Xe ions as obtained by different investigators, Western and USSR. They discuss the derivation of the formula for overcharge q cross-section by others. Fifteen references: e.g. Yu. N. Demkov, Uch, zap. LGU, No 146, ser. fiz. nauk, 8, 74, 1952; the authors, DAN SSSR, 98, 575, 1954; etc. (mostly Western references).

Institution: Karelo-Finnish State University

Submitted : May 30, 1955

*V.M. Kagan*

USSR/Physical Chemistry - Atom.

B-3

Abs Jour : Referat Zhur - Khimiya, No 1, 1958, 46

Author : V.M. Zakharova, Yu.M. Kagan.

Inst : -

Title : Study of Discharge Parameters and Character of Excitation  
of Ion Lines at Great Current Densities.

Orig Pub : Optika i Spektroskopija, 1956, 1, No 5, 627-635

Abstract : Sounding and optical measurements were carried out in dis-  
charges in Hg vapor (at pressures of  $1.2 \cdot 10^{-3}$  to  $1.2 \cdot 10^{-1}$  mm of mercury column) and in Ar, Kr and Xe (in the  
pressure range from 0.2 to 2.0 mm of mercury column) at  
current densities from 2 to 20 a per sq.cm. The ion part  
and the beginning of the electron part of the volt-ampere  
characteristic was used fro the measurement of the ion  
concentration. The electron concentration  $n_e$ , the tempera-  
ture of the electron gas  $T_e$  in inert gases does not change

APPROVED FOR RELEASE 08/10/2001 CIA RDP86-00513R000619910019-5"

Card 1/2

USSR/Physical Chemistry - Atom.

B-3

Abs Jour : Ref Zhur - Khimiya, No 1, 1958, 46

pressures of 1.0 and 1.75 mm of mercury column, while the  
same dependences obtained for Hg do not show any anomalies.  
The comparison of intensity curves of ion lines of Ar with  
curves of  $T_e$  and  $n_e$  under the same discharge conditions  
shows that the course of the intensity curves has a maxi-  
mum under the same conditions as the course of the concen-  
tration curves. It is postulated that the mechanism of  
ion excitation is stepwise.

"APPROVED FOR RELEASE: 08/10/2001

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APPROVED FOR RELEASE: 08/10/2001

CIA-RDP86-00513R000619910019-5"